Chlamydia trachomatis/Neisseria gonorrhoeae (Non-Culture) State of Michigan - Regional Laboratory Test Requisition

Date Received in Laboratory										Laboratory Sample #											
Michigan Department of Community Health-Bureau of Laboratories										•	Saginaw County Health Department										
3350 N. Martin Luther King Jr. Blvd.										1600 North Michigan											
PO Box 30035											Saginaw, Michigan 48602										
Lansing, Michigan 48909											Telephone: 989-758-3825										
Labora	atory Rec	ords: 51	7-335-80	59 Tech	nical Info	rmation:	517-335-	8067 Fa	x: 517-33	35-9871				Fa	ıx: 989	758-3	755				
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SPECIMEN INFORMATION																			
DATE COLLECTED (MM/DD/YYYY)								TIME COLLECTED											
18									19						O A	М		0	PM
SUBM	SUBMITTER'S SPECIMEN#																		
20																			
TEST	TEST REQUESTED																		
21	O C. trachomatis only (non-culture) O C. trachomatis and N. gonorrhoeae combo (non-culture)																		
SPEC	SPECIMEN SOURCE																		
22	O Cervix O Vagina O Urine O Urethra O Rectum (Lansing only) O Pharynx (Lansing only)							')											
REASON FOR TESTING - Check all areas that apply. (refer to definitions/explanations)																			
23	0.5	Sympto	ms			O His	story o	f STD ((<3 yea	3 years) O Age Recommended For Testing							ng		
23	O Infected Partner					O Partner Risk O Prenatal Visit O Retest										t			

Definitions/Explanations

Symptoms: Patient requesting examination due to symptoms, or, symptoms discovered upon

examination.

Infected Partner: Patient has known exposure to STD (self-reported or documented).

Partner Risk: Patient has multiple sex partners.

History of STD:

Patient has been diagnosed with a sexually transmitted disease within the last 3

vears.

Prenatal Visit: Patient examination is part of prenatal visit.

Age Recommended: CDC recommends annual screening of females ≤ 24.

Retest:

Patients diagnosed with chlamydia and gonorrhea should be retested approximately three (3) months after treatment, regardless of whether they believe that their sex partners were treated. If retesting at three months is not possible, clinicians should retest whenever that person next presents for medical care in the twelve months following initial treatment.

FP STD: This field is to be completed by sites supported by the Michigan Department of Community

Health to provide STD and/or Family Planning services. Completion of this field will assist

us in linking tests with the correct submitter site.

Zip Code: Patient zip code data is used to calculate screening rates in local jurisdictions and compare

them to infection. The resulting information can be used to better target resources and

testing.

Specimen Collection: Specimens must be collected using the appropriate collection kit as shown below.

Specimens received in the wrong collection kit will not be tested and reported as

"Unsatisfactory."

Specimen Source	Collection Kit
Endocervix, Urethra, Rectum, Pharynx	Aptima Unisex Swab
Urine	Aptima Urine Collection Kit
Vagina	Aptima Vaginal Swab

Rectal or Pharyngeal Swabs:

Limited testing of rectal and/or pharyngeal specimens is available only in the Lansing laboratory. This is not intended for population based screening: MDCH recommends the use of this test only for patients with symptoms or known exposure.